b.) Amendment to the Claims:

1. (Original) A method for quantitatively determining cholesterol in highdensity lipoprotein, which comprises:

reacting a sample with cholesterol esterase and cholesterol oxidase or cholesterol esterase, an oxidized coenzyme and cholesterol dehydrogenase in an aqueous medium comprising a bile acid derivative; and

measuring the formed hydrogen peroxide or a reduced coenzyme.

- 2. (Original) The method according to claim 1, wherein the aqueous medium further comprises albumin.
- 3. (Original) The method according to claim 1 or 2, wherein the cholesterol esterase is chemically modified cholesterol esterase.
- 4. (Original) The method according to claim 3, wherein the chemically modified cholesterol esterase is cholesterol esterase which is modified by a group selected from the group consisting of a group having poly(ethylene glycol) as a main component, a group having poly(propylene glycol) as a main component, a group having a copolymer of poly(propylene glycol) and poly(ethylene glycol), a group having a water-soluble polysaccharide, a sulfopropyl group, a sulfobutyl group, a polyurethane group and a group having a chelating function.

- 5. (Original) The method according to claim 3, wherein the chemically modified cholesterol esterase is cholesterol esterase which is modified by a group having poly(ethylene glycol) as a main component.
- 6. (Currently Amended) The method according to any one of claims 1 to 5 claim 4, wherein the bile acid derivative is a bile acid derivative having an anionic surface activity.
- 7. (Original) The method according to claim 6, wherein the bile acid derivative having an anionic surface activity is selected from the group consisting of cholic acid or a salt thereof, taurocholic acid or a salt thereof, glycocholic acid or a salt thereof, lithocholic acid or a salt thereof, deoxycholic acid or a salt thereof, chenodeoxycholic acid or a salt thereof, ursodeoxycholic acid or a salt thereof, 7-oxolithocholic acid or a salt thereof, 12-oxolithocholic acid or a salt thereof, 7-oxodeoxycholic acid or a salt thereof, hyocholic acid or a salt thereof, hyocholic acid or a salt thereof, hyocholic acid or a salt thereof,
- 8. (Currently Amended) The method according to any one of claims 1 to 5 claim 4, wherein the bile acid derivative is a bile acid derivative having a amphoteric surface activity.

9. (Original) The method according to claim 8, wherein the bile acid derivative having an amphoteric surface activity is a compound represented by the formula (1)

$$R^{1}$$
-CH₂-CH(R^{2})-CH₂-SO₃⁻ (I)

[wherein R^1 is a 3-(3-cholamidopropyl)dimethylammonio group and R^2 is a hydrogen atom or a hydroxyl group].

- 10. (Currently Amended) The method according to any one of claims 1 to 5 claim 4, wherein the bile acid derivative is a bile acid derivative having a nonionic surface activity.
- 11. (Original) The method according to claim 10, wherein the bile acid derivative having a nonionic surface activity is a compound represented by the formula (II)

(wherein X is a hydrogen atom or a hydroxyl group; R³ and R⁴ may be the same or different and each represents a substituted or unsubstituted alkyl group or a substituted or unsubstituted alkanoyl group) or a compound represented by the formula (III)

$$Z \xrightarrow{CH_3} Q \xrightarrow{O}_n W$$

$$X \xrightarrow{CH_3} Y$$

$$(111)$$

{wherein X, Y and Z may be the same or different and each represents a hydrogen atom, a hydroxyl group or an oxo (=O) group; Q is an oxygen atom or NH; W is a hydrogen atom, an alkyl group, an alkenyl group, an alkynyl group, a cycloalkyl group, a cycloalkenyl group, an alkanoyl group, an alkenoyl group, an alkynoyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted aminoalkyl group or a group represented by the formula (IV)

$$\begin{array}{c} \bullet \\ \\ N \\ H \\ m \end{array}$$

$$\begin{array}{c} CH_3 \\ \\ CH_3 \\ \\ Y' \end{array}$$

$$\begin{array}{c} CH_3 \\ \\ X' \end{array}$$

$$(IV)$$

- 12. (Original) A reagent for quantitatively determining cholesterol in highdensity lipoprotein, which comprises cholesterol esterase, cholesterol oxidase, a bile acid derivative and a reagent for quantitatively determining hydrogen peroxide.
- 13. (Original) A reagent for quantitatively determining cholesterol in highdensity lipoprotein, which comprises cholesterol esterase, cholesterol dehydrogenase, a bile acid derivative and an oxidized coenzyme.
- 14. (Original) The reagent according to claim 13, which further comprises a reagent for quantitatively determining a reduced coenzyme.
- 15. (Original) The reagent according to any one of claims 12 to 14, which further comprises albumin.
- 16. (Currently Amended) The reagent according to any one of claims 12 to 15 claim 15, wherein the cholesterol esterase is chemically modified cholesterol esterase.
- 17. (Original) The reagent according to claim 16, wherein the chemically modified cholesterol esterase is cholesterol esterase which is modified by a group selected from the group consisting of a group having poly(ethylene glycol) as a main component, a group having poly(propylene glycol) as a main component, a group having a copolymer of

poly(propylene glycol) and poly(ethylene glycol), a group having a water-soluble polysaccharide, a sulfopropyl group, a sulfobutyl group, a polyurethane group and a group having a chelating function.

- 18. (Original) The reagent according to claim 16, wherein the chemically modified cholesterol esterase is cholesterol esterase which is modified by a group having poly(ethylene glycol) as a main component.
- 19. (Currently Amended) The reagent according to any one of claims 12 to 18 claim 17, wherein the bile acid derivative is a bile acid derivative having an anionic surface activity.
- 20. (Original) The reagent according to claim 19, wherein the bile acid derivative having an anionic surface activity is selected from the group consisting of cholic acid or a salt thereof, taurocholic acid or a salt thereof, glycocholic acid or a salt thereof, lithocholic acid or a salt thereof, deoxycholic acid or a salt thereof, chenodeoxycholic acid or a salt thereof, ursodeoxycholic acid or a salt thereof, 7-oxolithocholic acid or a salt thereof, 12-oxochenodeoxycholic acid or a salt thereof, 7-oxodeoxycholic acid or a salt thereof, hyocholic acid or a salt thereof, hyocholic acid or a salt thereof, hyocholic acid or a salt thereof,

- 21. (Currently Amended) The reagent according to any one of claims 12 to 18 claim 17, wherein the bile acid derivative is a bile acid derivative having an amphoteric surface activity.
- 22. (Original) The reagent according to claim 21, wherein the bile acid derivative having an amphoteric surface activity is a compound represented by the formula (I)

$$R^{1}$$
-CH₂-CH(R^{2})-CH₂-SO₃ (I)

[wherein R^1 is a 3-(3-cholamidopropyl)dimethylammonio group and R^2 is a hydrogen atom or a hydroxyl group].

- 23. (Currently Amended) The reagent according to any one of claims 12-to 18 claim 17, wherein the bile acid derivative is a bile acid derivative having a nonionic surface activity.
- 24. (Original) The reagent according to claim 23, wherein the bile acid derivative having a nonionic surface activity is a compound represented by the formula (II)

(wherein X is a hydrogen atom or a hydroxyl group; R³ and R⁴ may be the same or different and each represents a substituted or unsubstituted alkyl group or a substituted or unsubstituted alkanoyl group) or a compound represented by the formula (III)

$$CH_3$$
 CH_3
 CH_3

(wherein X, Y and Z may be the same or different and each represents a hydrogen atom, a hydroxyl group or an oxo (=O) group; Q is an oxygen atom or NH; W is a hydrogen atom, an alkyl group, an alkenyl group, an alkynyl group, a cycloalkyl group, a cycloalkyl group, an alkanoyl group, an alkynoyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted aminoalkyl group or a group represented by the formula (IV)

$$\begin{array}{c}
\bullet \\
 \\
N \\
H
\end{array}
\right)_{m}
\begin{array}{c}
CH_{3} \\
CH_{3} \\
\end{array}$$

$$CH_{3}$$

$$CH_{3}$$

$$CH_{3}$$

$$V'$$

$$X'$$

- 25. (Original) A kit for quantitatively determining cholesterol in high-density lipoprotein, which comprises a first reagent comprising cholesterol esterase and a second reagent comprising cholesterol oxidase, wherein a bile acid derivative and a reagent for quantitatively determining hydrogen peroxide are comprised in either or both of the first reagent and/or the second reagent.
- 26. (Original) A kit for quantitatively determining cholesterol in high-density lipoprotein, which comprises a first reagent comprising a bile acid derivative and a second reagent comprising cholesterol esterase and cholesterol oxidase, wherein a reagent for quantitatively determining hydrogen peroxide is comprised in either or both of the first reagent and/or the second reagent.

- 27. (Original) A kit for quantitatively determining cholesterol in high-density lipoprotein, which comprises a first reagent comprising a reagent for quantitatively determining hydrogen peroxide a second reagent comprising cholesterol esterase and cholesterol oxidase wherein a bile acid derivative is comprised in either or both of the first reagent and/or the second reagent.
- 28. (Original) A kit for quantitatively determining cholesterol in highdensity lipoprotein, which comprises a first reagent cholesterol esterase and a second reagent comprising cholesterol dehydrogenase where a bile acid derivative and an oxidized coenzyme are comprised in either or both of the first reagent and/or the second reagent.
- 29. (Original) A kit for quantitatively determining cholesterol in high-density lipoprotein, which comprises a first reagent comprising a bile acid derivative and a second reagent comprising cholesterol esterase and cholesterol dehydrogenase wherein an oxidized coenzyme is comprised in either or both of the first reagent and/or the second reagent.
- 30. (Original) The kit according to claim 28 or 29, which further comprises a reagent for quantitatively determining a reduced coenzyme in either or both of the first reagent and/or the second reagent.

- 31. (Currently Amended) The kit according to any one of claims 25 to 30 claim 30, which further comprises albumin in either or both of the first reagent and/or the second reagent.
- 32. (Currently Amended) The kit according to any one of claims 25 to 31 claim 31, wherein the cholesterol esterase is chemically modified cholesterol esterase.
- 33. (Original) The kit according to claim 32, wherein the chemically modified cholesterol esterase is cholesterol esterase which is modified by a group selected from the group consisting of a group having poly(ethylene glycol) as a main component, a group having poly(propylene glycol) as a main component, a group having a copolymer of poly(propylene glycol) and poly(ethylene glycol), a group having a water-soluble polysaccharide, a sulfopropyl group, a sulfobutyl group, a polyurethane group and a group having a chelating function.
- 34. (Original) The kit according to claim 32, wherein the chemically modified cholesterol esterase is cholesterol esterase which is modified by a group having poly(ethylene glycol) as a main component.
- 35. (Currently Amended) The kit according to any one of claims 25 to 34 claim 33, wherein the bile acid derivative is a bile acid derivative having an anionic surface activity.

- 36. (Original) The kit according to claim 35, wherein the bile acid derivative having a anionic surface activity is selected from the group consisting of cholic acid or a salt thereof, taurocholic acid or a salt thereof, glycocholic acid or a salt thereof, lithocholic acid or a salt thereof, deoxycholic acid or a salt thereof, chenodeoxycholic acid or a salt thereof, ursodeoxycholic acid or a salt thereof, 7-oxolithocholic acid or a salt thereof, 12-oxochenodeoxycholic acid or a salt thereof, 7-oxodeoxycholic acid or a salt thereof, hyocholic acid or a salt thereof, hyocholic acid or a salt thereof,
- 37. (Currently Amended) The kit according to any one of claims 25 to 34 claim 33, wherein the bile acid derivative is a bile acid derivative having an amphoteric surface activity.
- 38. (Original) The kit according to claim 37, wherein the bile acid derivative having an amphoteric surface activity is a compound represented by the formula (I)

$$R^{1}$$
-CH₂-CH(R^{2})-CH₂-SO₃ (I)

[wherein R^1 is a 3-(3-cholamidopropyl)dimethylammonio group and R^2 is a hydrogen atom or a hydroxyl group].

39. (Currently Amended) The kit according to any one of claims 25 to 34 claim 33, wherein the bile acid derivative is a bile acid derivative having a nonionic a surface activity.

40. (Original) The kit according to claim 39, wherein the bile acid derivative having a nonionic surface activity is a compound represented by the formula (II)

$$\begin{array}{c}
CH_3 \\
OH \\
CH_3
\end{array}$$

$$\begin{array}{c}
CH_3 \\
H
\end{array}$$

$$\begin{array}{c}
R^4 \\
H
\end{array}$$

$$\begin{array}{c}
R^4 \\
H
\end{array}$$

$$\begin{array}{c}
R^4 \\
H
\end{array}$$

(wherein X is a hydrogen atom or a hydroxyl group; R³ and R⁴ may be the same or different and each represents a substituted or unsubstituted alkyl group or a substituted or unsubstituted alkanoyl group) or a compound represented by the formula (III)

$$Z$$
 CH_3
 CH_3
 CH_3
 CH_3
 $(IIII)$

{wherein X, Y and Z may be the same or different and each represents a hydrogen atom, a hydroxyl group or an oxo (=O) group; Q is an oxygen atom or NH; W is

a hydrogen atom, an alkyl group, an alkenyl group, an alkynyl group, a cycloalkyl group, a cycloalkenyl group, an alkanoyl group, an alkenoyl group, an alkynoyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted aminoalkyl group or a group represented by the formula (IV)

$$\begin{array}{c|c} \bullet & \bullet & \bullet \\ & N \\ & N \\ & H \\ & m \end{array} \begin{array}{c} CH_3 \\ & CH_3 \\ & \\ & Y' \end{array} \hspace{1cm} (IV)$$

41. (Original) A compound represented by the formula (III)

$$CH_3$$
 CH_3
 CH_3

{wherein X, Y and Z may be the same or different and each represents a hydrogen atom, a hydroxyl group or an oxo (=O) group; Q is an oxygen atom or NH; W is a hydrogen atom, an alkyl group, an alkenyl group, an alkynyl group, a cycloalkyl group, a cycloalkyl group, an alkenyl group, an alkynoyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted aminoalkyl group or a group represented by the formula (IV)

$$\begin{array}{c} \bullet \\ \\ N \\ H \\ m \end{array} \begin{array}{c} CH_3 \\ CH_3 \\ \\ Y' \end{array} \begin{array}{c} CH_3 \\ \\ X' \end{array} \tag{IV}$$

42. (Original) A process for producing a compound represented by the formula (III)

$$CH_3$$
 CH_3
 CH_3

(wherein X, Y and Z may be the same or different and each represents a hydrogen atom, a hydroxyl group or an oxo (=O) group; Q is an oxygen atom or NH; W is a hydrogen atom, an alkyl group, an alkenyl group, an alkynyl group, a cycloalkyl group, a cycloalkenyl group, an alkanoyl group, an alkenyl group, an alkynoyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted aminoalkyl group or a group represented by the formula (IV)

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integer of 3 to 400}, which comprises: reacting a compound represented by the formula (V)

$$CH_3$$
 CH_3
 CH_3
 CH_3
 CH_3
 (V)

[wherein X, Y and Z may be the same or different and each represents a hydrogen atom, a hydroxyl group or an oxo (=O) group] with a compound represented by the formula (VI)

$$HO$$
 O
 W'
 (VI)

(wherein W' is a hydrogen atom, an alkyl group, an alkenyl group, an alkenyl group, a cycloalkyl group, a cycloalkenyl group, an alkanoyl group, an alkenoyl group, an alkynoyl group or a substituted or unsubstituted aryl group; and n is an integer of 3 to 400) or with a compound represented by the formula (VII)

$$H_2N$$
 (VII)

(wherein T is a substituted or unsubstituted aminoalkyl group; and n is an integer of 3 to 400) in the presence of a condensing agent.